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**Transport
diagnostics for the
lowermost
stratosphere**

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Observationally derived transport diagnostics for the lowermost stratosphere and their application to the GMI chemistry and transport model

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Transport from the surface to the lowermost stratosphere can occur on timescales of a few months or less, making it possible for short-lived tropospheric pollutants to influence stratospheric composition and chemistry. Models used to study this influence must demonstrate the credibility of their chemistry and transport in the upper troposphere and lower stratosphere (UT/LS). Data sets from satellite and aircraft instruments measuring CO, O₃, N₂O, and CO₂ in the UT/LS are used to create a suite of diagnostics of the seasonally-varying transport into and within the lowermost stratosphere, and of the coupling between the troposphere and stratosphere in the extratropics. The diagnostics are used to evaluate a version of the Global Modeling Initiative (GMI) Chemistry and Transport Model that uses a combined tropospheric and stratospheric chemical mechanism and meteorological fields from the GEOS-4 general circulation model. The diagnostics derived from N₂O and O₃ show that the model lowermost stratosphere (LMS) has realistic input from the overlying high latitude stratosphere in all seasons. Diagnostics for the LMS show two distinct layers. The upper layer (~350 K–380 K) has a strong annual cycle in its composition, while the lower layer, just above the tropopause, shows no seasonal variation in the degree of tropospheric coupling or composition. The GMI CTM agrees closely with the observations in both layers and is realistically coupled to the UT in all seasons. This study demonstrates the credibility of the GMI CTM for the study of the impact of tropospheric emissions on the stratosphere.

1 Introduction

Tropospheric pollutants can impact the composition and chemistry of the lower stratosphere. Using CO measurements from the Microwave Limb Sounder (MLS) onboard the NASA AURA satellite, Schoeberl et al. (2006) identified a “tape recorder” of CO in the tropical upper troposphere forced by seasonal variations in biomass burning.

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With a lifetime of several months, CO is sufficiently long-lived for the tape recorder to be observed at 20 km in the tropical lower stratosphere (LS). Short-lived brominated tropospheric source gases such as bromoform (CHBr₃) may affect lower stratospheric chemistry. They are typically ignored in the calculation of stratospheric Br_y because of their short lifetimes, but Ko et al. (1997) used a 2D model to illustrate that Br_y produced from short-lived species in the upper troposphere may contribute to stratospheric halogen loading. Based on a comparison of observed BrO with model calculations, Salawitch et al. (2005) suggested the existence of a 4–8 ppt Br_y source in the tropical upper troposphere that is rapidly transported to the lowermost stratosphere (LMS), where it increases ozone loss. These studies provide examples of the potential for tropospheric pollutants with lifetimes of a few months to change the composition and chemistry of the LMS.

There are two major pathways for entry to the stratosphere. Convection in the tropics brings tropospheric air up to the base of the tropical transition layer (TTL), which begins at ~13 km (~350 K) (Schoeberl et al., 2006). Net heating rates become positive in the TTL and ascent by the Brewer-Dobson circulation slowly lifts air up to and across the tropical tropopause (17–18 km) and into the lower stratosphere. A second pathway involves entry of tropospheric air into the extratropical LMS by quasi-horizontal transport out of the TTL. This pathway is aided by monsoon anticyclones in the summer hemisphere (Chen, 1995), with poleward transport of tropospheric air on the west side and equatorward transport of stratospheric air on the east side of the monsoonal circulation. Asia and Mexico are the most active areas for monsoon transport in the northern summer.

The extratropical lowermost stratosphere (LMS) is defined as the region between the extratropical tropopause, where isentropes connect the stratosphere and troposphere, and roughly 380 K (~100 hPa), where all isentropes are in the stratosphere (Fig. 1). This is the stratospheric part of the middleworld defined by Hoskins (1991). The LMS is composed of air of both tropospheric and stratospheric origin, and their relative fractions vary with season. In winter, large potential vorticity (PV) gradients

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near the subtropical jet form an elastic, nearly impermeable barrier to isentropic transport much like that found at the edge of the polar vortex. Subtropical PV gradients are weaker in summer and evanescent waves from monsoonal circulations allow considerable stratosphere-troposphere exchange (STE) between the tropical upper troposphere (UT) and the LMS (Chen, 1995). Diagnostics of LMS transport, therefore, evaluate the integrated effects of the Brewer-Dobson (stratospheric) and monsoon (tropospheric) circulations.

There is agreement in the chemistry-climate model community on the need for diagnostics in the UT/LMS. There are numerous trace gas measurements in the UT/LMS whose seasonal and spatial variations make them suitable for the development of transport diagnostics. Trace gases with large or seasonally-varying gradients across the tropopause are excellent indicators of the strength and timing of the dynamical processes affecting the UT/LMS. For example, Ray et al. (1999) used CO₂ and CFC-11 profiles to demonstrate that LMS composition changes from primarily stratospheric to tropospheric air between spring and fall. Hoor et al. (2002) noted differences in the CO-O₃ correlation between winter and summer that indicated increased influence of tropospheric air in the lowermost stratosphere in July. The observed composition changes represent the shifting balance between forcings with different seasonal maxima, i.e., the downward branch of the Brewer-Dobson circulation and the subtropical summer monsoons. The analysis of these observations may lead to useful diagnostics that show the net effect of the forcings.

In this paper, we present diagnostics of UT/LMS transport and composition derived from satellite and aircraft measurements of CO, CO₂, N₂O, and O₃. Dynamical processes relevant to the UT/LMS include the ascent of tropical air from the surface to the stratosphere, the seasonally-varying transport from the tropical UT to the LMS, and the coupling between the extratropical LMS and the UT. We also report on a new version of the Global Modeling Initiative (GMI) chemistry and transport model (CTM) that uses meteorological fields from the GEOS-4 general circulation model (GCM) (Bloom et al., 2005) and has a chemical mechanism that includes tropospheric and stratospheric

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chemical reactions. The diagnostics are applied to model results to establish the credibility of the meteorological representation of the UT/LMS. The evaluations demonstrate the viability of using this CTM to predict the impact of the tropospheric emissions on lower stratospheric composition.

2 Model and simulation descriptions

The GMI CTM used in this study is similar to the version described in Douglass et al. (2004) and references therein. The CTM uses a flux form semi-Lagrangian numerical transport scheme (Lin and Rood, 1996). The version of the model used here, referred to as the 'GMI Combo CTM', has a chemical mechanism that combines the stratospheric mechanism described in Douglass et al. (2004) with a modified version of the tropospheric mechanism originating in the Harvard GEOS-CHEM model (Bey et al., 2001). This combined mechanism contains 113 chemically active species, 315 chemical reactions and 78 photolytic processes. A new version of the Fast-J2 photochemical solver (Bian and Prather, 2002) called Fast-JX includes more bins for the calculation of stratospheric photolysis rates. Photolysis frequencies are computed using the Fast-JX radiative transfer algorithm, which combines the Fast-J tropospheric photolysis scheme described in Wild et al. (2000) with the Fast-J2 stratospheric photolysis scheme of Bian and Prather (2002) (M. Prather, personal communication, 2005). The scheme treats both Rayleigh scattering as well as Mie scattering by clouds and aerosol. The SMVGEAR II solver uses a 30-min timestep for the chemistry calculation, which is more accurate near the terminator than the 1-hr timestep used in previous studies. Performance metrics of SMVGEAR II with respect to other solvers in the GMI stratospheric CTM can be found in Rotman et al. (2001). Model representation of physical processes in the troposphere, such as convection, wet scavenging, and dry deposition, are described in B. Duncan et al. ("Model study of the cross-tropopause transport of biomass burning pollution", submitted manuscript).

Long-lived source gases, such as N_2O , CH_4 , and the halocarbons, are forced at

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the two lowest levels with a mixing ratio boundary condition updated monthly, based on the A2 scenario (WMO, 2002). A climatology is used to specify the stratospheric distribution of water vapor each month. The GMI model calculates a change in water from the specified trend in methane and adds this to the climatology at each step. CO₂ is forced at the two lowest levels with a mixing ratio boundary condition using a time series derived from global surface observations (Conway et al., 1994) from a 17-yr period. The CO₂ boundary condition has 10°-wide latitude bins with no longitudinal variability and is updated monthly. Model CO sources include emissions from biomass burning, fossil fuel consumption, biofuel use, lightning, and biogenics. They are treated as fluxes rather than mole fraction boundary conditions and are described in Duncan et al., submitted manuscript. This mechanism lacks a high altitude loss for CO₂, a source of CO, and thus the model CO is biased low in the stratosphere.

The CTM simulation evaluated here was run with meteorological fields from a 5-yr integration of the GEOS-4.0.2 GCM (Bloom et al., 2005). This integration was forced by observed sea surface temperatures for the period 1994–1998. The native resolution of the meteorological fields is 2° latitude by 2.5° longitude and 55 vertical levels with a lid at 0.015 hPa. Resolution in the UT/LMS is 1 km or less. To shorten the CTM integration time, the 24 levels above 10 hPa were mapped to 11 levels using the method of Lin (2004); the top level is unchanged. The resulting CTM grid is 2° latitude × 2.5° longitude by 42 levels. A study of the effects of resolution and lid height on CTM transport characteristics determined that transport in the lower stratosphere and below is negligibly impacted by reduced resolution above 10 hPa (Strahan and Polansky, 2006).

3 Transport diagnostics in the upper troposphere and lower stratosphere

Evaluation of a stratospheric CTM simulation integrated with meteorological fields from a GCM essentially evaluates extratropical wave driving in the GCM, the driver of the stratospheric circulation. A GCM that correctly simulate the generation, propagation, and dissipation of Rossby waves may correctly represent the global scale aspects of

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transport and STE (Holton et al., 1995). However, composition of the lowermost stratosphere depends on the interaction of upper tropospheric processes with the Brewer-Dobson circulation. Thus, the diagnostics presented here evaluate the integrated effect of stratospheric and tropospheric processes on the UT/LMS.

5 3.1 Transport up to and through the tropical tropopause

Changes in the amplitude and phase of the CO₂ cycle observed at points 1 through 4 in Fig. 1 provide the basis for a series of transport diagnostics. Boering et al. (1996) (hereinafter referred to as B96) used ground-based CO₂ measurements (Conway et al., 1994) along with NASA ER-2 aircraft data to estimate transport rates from the surface to the tropical tropopause, and from the tropical LS and to the midlatitude LS. Figure 2 uses the B96 analysis to evaluate model transport from the surface to the tropical UT and beyond. B96 showed that the seasonal cycle at stratospheric entry (~390 K) could be represented by the average of the surface cycles measured at Mauna Loa (19° N) and Samoa (14° S) time lagged by 2 months (Fig. 2a); this is the transport between points 1 and 2 in Fig. 1. The solid black line is the average of the model surface forcing at Mauna Loa and Samoa, and the dashed line is that cycle lagged by 2 months. The model cycle at the tropical tropopause (red) is correctly lagged by 2 months but has an amplitude ~1 ppm less than observed, apparently a result of the surface minimum arriving with ~1 ppm higher CO₂ (in fall). From the tropical tropopause, B96 found that the cycle was transported upward to ~435 K in 3-4 months, diminished in amplitude by only 20%; this is the transport between points 2 and 3 in Fig. 1. If the model cycle at the tropopause (Fig. 2b, red) showed the same behavior, it would produce a cycle at 435 K shown by the dashed blue line in Fig. 2b. The actual model cycle at 435 K is shown by the solid blue line, which suggests that ascent is a little too rapid (the lag is 2.5 months) and that attenuation or horizontal exchange in the LS is too great, with ~40% loss of amplitude instead of only 20%. The loss of amplitude occurs during the ascent of the cycle maximum to 435 K in fall, suggesting that the Brewer-Dobson circulation may be too strong in this season. Schoeberl et al. (2006) also assessed tropical ascent

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and horizontal mixing in this model from 14–20 km by comparison with the MLS CO “tape recorder”. They found that the model agreed well with the observed tilt of the CO signal (ascent rate) and the height at which the signal faded (horizontal mixing convolved with CO lifetime). While the tape recorder evaluation was only qualitative, it suggests that ascent and mixing throughout the year are in the right ballpark; the CO₂ phase and attenuation comparison is a stricter test of these processes and shows good agreement from midwinter to summer, but suggests excess strength in fall.

B96 also examined poleward horizontal transport to the LMS; this is transport between points 2 and 4 in Fig. 1. Their CO₂ analysis used potential temperature and co-located N₂O mixing ratios to create a vertical coordinate that was effectively tied to the extratropical tropopause (see Hoor et al., (2004), hereinafter referred to as H2004). They found that the CO₂ cycle observed at the tropical tropopause could also be found near ~38° N, undiminished, with no more than a 1-month time lag. Figure 2c shows the model CO₂ cycles in the tropics and from 36–40° N, sorted by N₂O=305–310 ppb and theta=380–400 K, just as in the B96 analysis. The near perfect match of the phase and amplitude of these cycles suggests realistic poleward transport in the 380 K region of the stratosphere. Notice that there is no lag in summer and fall, but roughly a 1 month lag in winter and spring. This is consistent with transport variations in the LMS described by Chen (1995) and Dunkerton (1995), who showed stronger exchange between the tropics and midlatitudes occurring in summer.

3.2 The seasonally-varying composition of the lowermost stratosphere

Several analyses have shown that the composition of the LMS varies considerably between winter and summer (e.g., Ray et al., (1999); Hoor et al., (2002); and H2004). In winter and spring, the LMS has a stratospheric character due to strong downward motion at mid and high latitudes while large PV gradients across the subtropical jet restrict horizontal transport. In summer, when downward advection by the Brewer-Dobson circulation is weak, the LMS has greater tropospheric character due to weak subtropical PV gradients which allow enhanced horizontal transport from the tropical

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UT to the LMS by the monsoonal anticyclones (Chen, 1995).

Ozone, N₂O, and CO are useful as tracers of the origin of air because of their gradients across the tropopause. N₂O and O₃ have opposite source regions, the troposphere and stratosphere, respectively, and are long-lived in the LMS. CO has a tropospheric source and a lifetime of several months in the LMS; its tropospheric sources have a strong semi-annual cycle due to seasonal biomass burning from both hemispheres (Schoeberl et al., 2006).

Several datasets will be used to examine seasonal variations in lower stratospheric composition. The Microwave Limb Sounder (MLS) on the NASA Aura satellite provides global measurements for O₃ (215 hPa and above) and N₂O (100 hPa and above). MLS N₂O at 100–68 hPa shows occasional high bias estimated to be generally less than 10%; MLS O₃ has a reported bias of 1% in the stratosphere and 10% or less in the tropical UT (Livesey et al., 2005). N₂O data from eight SPURT aircraft campaigns are available in the extratropical LMS in all seasons (Engel et al., 2006). Seasonal mean analyses of ER-2 N₂O and O₃ data [Strahan et al., 1999; Strahan, 1999] provide profiles and latitudinal gradients from 360–500 K. We use a dynamical definition of the tropopause, 2 PVU, where 1 PVU = 10⁻⁶ m² K s⁻¹ kg⁻¹, to define the lower boundary of the LMS.

3.2.1 Lower stratospheric N₂O

Through downward transport by the Brewer-Dobson circulation, the lower stratosphere is a source of air for the LMS, thus its composition is relevant. Low N₂O descends to the LMS in winter and spring, while horizontal poleward transport in summer brings higher (tropospheric) mixing ratios. Figure 3 shows seasonal contoured pdfs of model N₂O, the most probable profile from pdfs of MLS N₂O (white lines), and seasonal mean N₂O profiles from ER-2 and SPURT measurements (white and red points, respectively). The contoured pdfs show the most probable values and the range of model variability found. The most probable values in winter represent the vortex mean, while the large variability comes from profiles outside the vortex. (The spring ER-2 data are omitted

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because they are from a year with an unusually late vortex breakup (Coy et al., 1997). The only disagreement with the data is found in summer and fall above 450 K, where the model is up to 15% too high. Although the strength of descent and horizontal mixing cannot be judged independently, the excellent model agreement throughout most of the year at levels as low as 320 K suggests a very good balance of vertical and horizontal transport in the polar lower stratosphere. The SPURT measurements are higher than the model in the LMS in winter, but this could be due to weaker descent in the warm Arctic winters in the years of the SPURT campaign (2002 and 2003). The model Arctic temperatures are near or below the real climatological mean.

Variability is an indicator of the presence or absence of transport processes. In winter, large tracer gradients across the vortex edge coupled with wave-driven vortex wobble create high latitude pdfs with large variability. Strong horizontal mixing in spring causes breakdown of the vortex and homogenization of the extratropics. With very weak wave-driving until mid-fall, the well-mixed extratropics produce pdfs with very low variability in the summer months. Figure 4 demonstrates the realistic seasonal variability of high latitude transport and mixing processes in the GMI model. In February, excellent agreement between MLS and the model most probable profiles inside and outside the vortex, and the variability in both regions, shows that the model vortex is well isolated. The model has more points between the two profiles, indicating slightly more mixing across the vortex edge. In summer, the MLS data and the model both show low variability characteristic of a well-mixed atmosphere with little wave-driving. Close agreement with MLS variability is also seen in the months not shown.

Figures 3 and 4 show that the model lower stratosphere provides realistic input for the LMS in all seasons. To examine transport characteristics and composition in the LMS, we look at SPURT N₂O in the dynamical coordinate system of equivalent latitude and potential temperature. The use of equivalent latitude, which maps each measurement onto latitude according to its potential vorticity, removes variability caused by reversible wave transport [Butchart and Remsberg, 1986; Nash et al., 1996]. H2004 and Hegglin et al. [2006] used this coordinate system to show that the isopleths of CO,

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N_2O , and O_3 follow PV contours rather than those of potential temperature in the LMS ($<360\text{ K}$) in all seasons. Figure 5 shows the model and SPURT N_2O in the spring and fall; the dashed lines are PV contours (2, 4, and 6 PVU). Note that the PV contours cross isentropes, or equivalently, there are PV gradients on the isentropic surfaces.

5 The PV gradients hinder horizontal transport, allowing time for diabatic cooling in the middleworld to move tracer isopleths downward as they are transported poleward. Because the model N_2O isopleths, like the data, clearly follow the dynamical tropopause and PV contours, this suggests the model has a reasonable balance between horizontal transport and diabatic cooling in the LMS. In all seasons except winter, model N_2O mixing ratios are always within 2% of SPURT values. As noted before, SPURT values are probably high in winter due to weak descent in the observation years.

Figure 5 also contrasts the composition differences between spring and fall. In spring, downward advection by the Brewer-Dobson circulation is at a maximum, resulting in relatively low N_2O , and in fall, the downward circulation is weak and the influence of horizontal transport from the tropics is greatest (higher N_2O), consistent with the Hegglin et al. (2006) analysis of transport using SPURT N_2O . The bottom panels show the ratio of fall to spring mixing ratios. The ratio is proportional to the seasonally changing tropospheric fraction of air in the LMS and is very well represented by the model as a function of both height and latitude.

20 3.2.2 Ozone

Ozone is also an excellent lower stratospheric transport tracer. The availability of global MLS O_3 measurements down to 215 hPa allows evaluation of the entire LMS and the tropical UT (the middleworld). Ozone in the middleworld is controlled largely by transport, although O_3 production and loss are sensitive to local NO_x . Figure 6 shows the annual cycle of O_3 from the tropics to high latitudes, from 350 K–420 K using MLS data (black). Model O_3 data are overlaid in red. (ER-2 data are not used here because they are only available as seasonal means.) Overall, the model shows exceptional agreement with the observed annual cycles and their variability. Model ozone in the

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tropics is ~ 50 ppb too low at all levels but shows the right seasonal variation, including the summer maximum seen in the MLS data. The low model O_3 could indicate an insufficient source of O_3 from lightning-produced NO_x in the tropical UT, or insufficient exchange with the midlatitudes, a source of higher O_3 . This discrepancy is so small that as a percentage it appears insignificant at higher levels ($<10\%$). The model high latitude O_3 shows near perfect agreement in all seasons at all levels shown. Model midlatitude O_3 looks good in summer and fall but is low in winter. This cannot easily be explained by horizontal transport problems. If the vortex were too isolated, this would act to reduce the amount of high O_3 mixed into the midlatitudes; however, Fig. 4 shows that the vortex is not overly isolated. And tropical O_3 is low, indicating, if anything, too little exchange with midlatitudes. This leaves insufficient midlatitude descent as a possible cause for the low O_3 in winter. Overall, the agreement between observed and model O_3 in the middleworld and LS shown in Fig. 6 is quite close when put in the context of the large range of observed mixing ratios here. Because variability and seasonal composition are controlled by transport, the O_3 and N_2O diagnostics demonstrate credible model transport throughout the year in the extratropical lower stratosphere from 320 K–500 K.

3.3 Stratosphere-troposphere coupling at the extratropical tropopause

3.3.1 CO and O_3

CO and O_3 have different source regions and hence different correlations in the troposphere and stratosphere. Near the extratropical tropopause, these opposite correlations are connected by lines with CO and O_3 mixing ratios intermediate between their tropospheric and stratospheric values (Pan et al., 2004; H2004). H2004 assessed the degree of coupling between the stratosphere and troposphere by measuring the thickness of the mixed layer near the tropopause. Using the dynamical tropopause (2 PVU) as a reference, they found that this mixed layer began just below the tropopause and extended to ~ 25 K above it, and that the thickness of this layer was the same in all sea-

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sons and latitudes in the extratropics. They also found that the thermal tropopause was usually slightly above the 2 PVU tropopause. On an absolute scale, the 350 K surface was almost always above the mixed layer during the SPURT campaigns at latitudes 50° N or higher. These results are consistent with the tracer studies of Chen(1995), who reported that vigorous exchange between the stratosphere and troposphere occurred year-round at 330 K and below due to breaking synoptic scale baroclinic disturbances. This is in contrast to their “middle” middleworld results (350 K and above), where seasonal mixing across the subtropical jet was found to be enhanced by the monsoon anticyclones in summer and prohibited by large PV gradients in winter.

We diagnose the model’s strat-trop interaction region by the thickness of its mixed layer and location with respect to the dynamical tropopause. This method was used on aircraft observations by Hoor et al. (2002). Figure 7 shows model CO and O₃ where heights with respect to the tropopause are color coded. At >30 K above the tropopause (green), CO is near its minimum (stratospheric) value, while <−10 K below the tropopause (blue), O₃ is near its minimum (tropospheric) value. The red points form mixing lines between the two regions and range from −10 K below to 30 K above the tropopause. The mixing lines are broadest in spring due to large variability in the upper tropospheric and lower stratospheric reservoir regions, indicating that different air masses (e.g. vortex and nonvortex) are involved in mixing. The mixing lines are most compact in fall when the reservoir regions are more homogeneous. We find that the thickness of the model mixed layer is nearly constant with latitude and season and the mixed layer extends no more than 30 K above the dynamical tropopause, just as in the SPURT data analysis (H2004). This suggests that the model provides reasonable forcing by synoptic scale disturbances in the UT throughout the year that couple the model’s troposphere with the lowest layers of the stratosphere.

3.3.2 CO₂

The CO₂ seasonal cycle phase and amplitude at the extratropical tropopause not only demonstrate the coupling between troposphere and stratosphere, but can be used to

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diagnose the influence of air arriving via the tropical stratosphere at levels above the coupling. Figure 8 shows the model CO₂ cycle 40°–60° N at several levels above and below the 2 PVU tropopause. The observed cycles in the UT and at the tropopause both have a spring maximum and a large amplitude (blue), but above that there are significant differences in phase and amplitude (Nakazawa et al., 1991; H2004). At +20 K (green) the amplitude is much lower than below and the maximum occurs more than a month later; these features suggest a mixture of influences from above and below, consistent with the mixed region diagnosed by the CO–O₃ scatterplots. At +40 K and above (orange and red), a purely stratospheric cycle appears, with low amplitude and a maximum shifted to late summer. As has been demonstrated by analysis of observations in H2004, B96, and Strahan et al. (1998), this cycle arrives via ascent through the tropical tropopause, followed by meridional poleward transport. The seasonal maximum that occurs in May in the northern mid-high latitude lower troposphere requires ~3 months to travel to the midlatitude LS via the tropical tropopause. A key feature of this diagnostic, shown clearly in Fig. 9 of H2004, is the presence of a reversed vertical gradient in late summer, with low CO₂ in the UT and high CO₂ in the LS (near day 250). In all other seasons CO₂ decreases with height. The timing and magnitude of the model gradient reversal is in good agreement with the H2004 analysis that showed a 2–3 ppm difference between the UT and 40–60 K above the tropopause in August. A model run with convection turned off failed to produce the low CO₂ in the UT in summer and the reversed gradient, demonstrating the importance of convection to correctly simulate composition near the tropopause. The GMI model shows a UT amplitude of 5–6 ppm, in excellent agreement with Fig. 9 of H2004.

4 Diagnostics summary and GMI Combo Model Credibility

Table 1 summarizes the suite of transport diagnostics presented in this paper. They have been applied to the GMI GEOS4-GCM Combo CTM to evaluate transport from the tropical upper troposphere to the lower stratosphere, composition of the lowermost

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stratosphere, and coupling between the troposphere and stratosphere. Most comparisons with data showed excellent agreement, but a few discrepancies were found in the analysis of tropical transport. Ascent from tropical tropopause to 435 K appeared to be a little too rapid in summer and the CO₂ cycle amplitude was attenuated too much as it ascended, suggesting too strong mixing with the midlatitudes. This is consistent with summer polar N₂O profiles being too high (too tropospheric). The Brewer-Dobson circulation in the GEOS4-GCM appears to be too strong at these altitudes in the warm seasons.

The observationally-derived diagnostics indicate two distinct regions within the lowermost stratosphere. The upper region, ~350 K–380 K, has seasonally varying composition, while the lower region, a ~30 K thick layer above the tropopause, shows uniform composition (excluding CO₂) and thickness year-round. In this lower region, the bulk of the comparisons demonstrate that the GMI Combo CTM has composition and transport characteristics that mirror the observations. The thickness of the mixed layer separating the troposphere and stratosphere (<350 K) agrees closely year-round with the SPURT data, and the phase and amplitude change of the CO₂ cycle across the tropopause is in agreement with the Nakazawa et al. (1991) and H2004 results. This suggests that the UT wave disturbances involved in creating the mixed layer (Chen, 1995) operate realistically year-round. SPURT data analyses have shown that transport of long-lived species (N₂O, O₃, and CO) follow isopleths of PV rather than potential temperature. Model tracer transport also follows PV isopleths. Downward transport out of the LMS (stratosphere-to-troposphere flux) was not been evaluated here but was been previously estimated for meteorological fields from the same GCM (Olsen et al., 2004). They compared ozone fluxes with empirically derived mass fluxes (Olsen et al., 2003) and found the absolute values and their seasonal variations were reasonable.

There is also good agreement in the middle and upper LMS (350–380 K) where there is a clear annual cycle in the composition. The annual cycle arises from the seasonally-varying influence of the stratosphere (strong downward motion in winter) and troposphere (horizontal transport of tropical air via monsoon anticyclones in sum-

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mer). Realistic model transport in this region is supported by O₃ seasonal cycles from the tropics to high latitudes which match extremely well with MLS observations for 2005, and by the ratio of fall/spring N₂O, which shows the seasonal influence from the troposphere and matches the SPURT data very closely. Middleworld composition depends on the composition of stratospheric air descending from above, and the close agreement of high latitude N₂O profiles with MLS, SPURT, and ER-2 data suggests realistic input from the overlying lower stratosphere in all seasons.

Model evaluation in the lower stratosphere is essential because a credible lower stratosphere is a prerequisite for a credible middleworld. Realistic transport into and within the lowermost stratosphere depends on Brewer-Dobson circulation in the meteorological fields used in a CTM. In the analysis of a similar CTM using GEOS4-GCM fields, Strahan and Polansky (2006) found that realistic lower stratospheric transport barriers (e.g., those near the polar vortex and in the subtropics) were essential for obtaining credible O₃ and CH₄ distributions. Meteorological fields known to have good barriers (e.g., the GEOS4-GCM) still require a CTM horizontal resolution of at least 2° × 2.5° in order to correctly produce the barriers in the CTM. Observationally-derived transport diagnostics used in Strahan and Polansky (2006) as well as several previous GMI studies are recommended for evaluation of stratospheric transport (Douglass et al., 1999; Strahan and Douglass, 2004).

The realistic seasonal cycle of transport into and within the GMI lowermost stratosphere shown here demonstrates the utility of this model for studies of the influence of tropospheric composition on the lower stratosphere. The model shows a reasonable transport time from the surface to the tropical upper troposphere, and in summer and fall the model transports tropical UT air into the lowermost stratosphere (~350 K–380 K). This demonstrates the feasibility of using this model to study the effects of tropospheric pollutants with a life time of even a few months on the composition, chemistry, and radiative properties of the LMS. The seasonally varying O₃ composition matches extremely well with observations, which supports the use of this model in studies involving perturbations to O₃. This model can be credibly used to study problems such as the

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impact of large emissions from fires on the stratosphere and the impact of short-lived halogenated species on ozone loss in the lowermost stratosphere.

Acknowledgements. This work is supported by the NASA Model Analysis and Prediction Program. We thank J. Rodriguez, Project Scientist of the Global Modeling Initiative for scientific support and E. Nielsen for producing the GEOS-4-GCM meteorological fields. We also thank N. Livesey and L. Froidevaux for use of the MLS version 1.5 ozone data.

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Table 1. Transport diagnostics for the upper troposphere and lowermost stratosphere.

Region	Feature	Reference
Surface to UT	CO ₂ cycle phase and amplitude at tropical stratospheric entry (~380 K)	Boering et al. (1996)
Troposphere to Stratosphere (vertical)	CO ₂ cycle phase and amplitude at 435 K in tropics	Boering et al. (1996)
Troposphere to Stratosphere (isentropic)	CO ₂ cycle phase and amplitude at ~380 K, 40° N	Boering et al. (1996)
Composition and Coupling in the LMS, Transport within the LMS	O ₃ seasonal cycle amplitude and variability, tropics to high latitudes, 350 K–420 K	This study.
	N ₂ O Seasonal Profiles, 70°–88° N, 320–500 K	This study.
	Ratio of Fall/Spring N ₂ O, 320–380 K (change in tropospheric fraction of LMS air)	This study.
	CO, O ₃ , and/or N ₂ O isopleths following the tropopause	Hoor et al. (2004)
	Consistent thickness of mixed layer at the extratropical tropopause	Hoor et al. (2004)
	Change in CO ₂ cycle amplitude from the UT to the LS	Nakazawa et al. (1991), Hoor et al. (2004)

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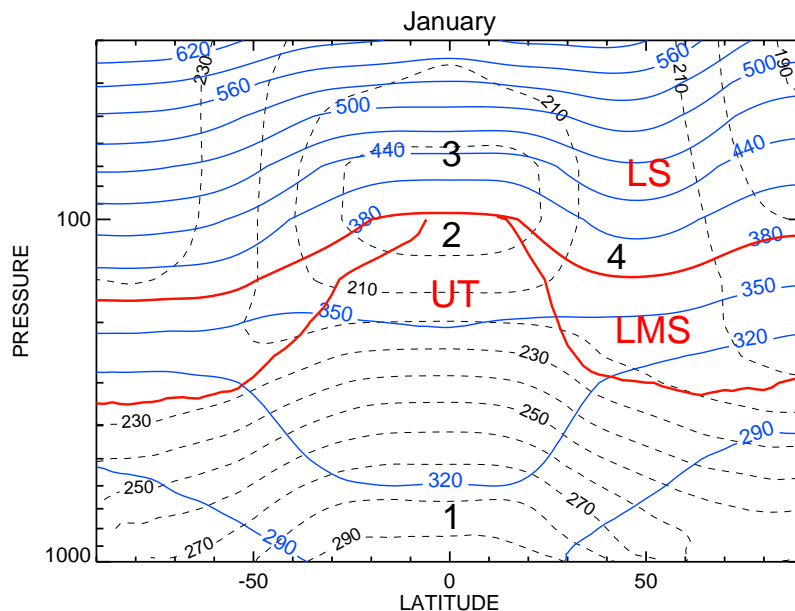


Fig. 1. Schematic diagram of the lowermost stratosphere derived from zonal monthly mean GEOS-4 meteorological analyses from January, 2005. Temperature contours are black (dashed), potential temperature contours are blue (solid), and the lowermost stratosphere, defined by the 380 K potential temperature surface and the 2 PVU surface, is outlined in red. Transport between the numbered regions is discussed in the text.

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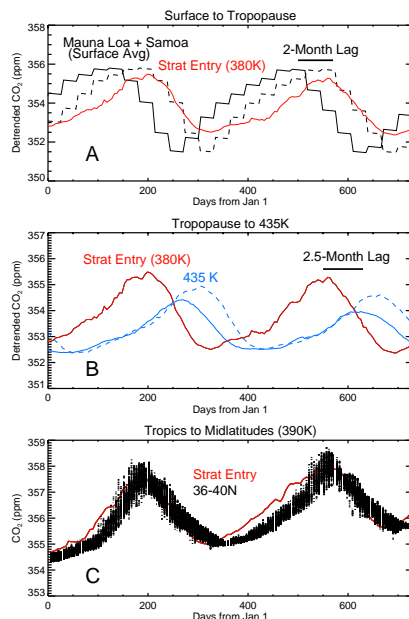


Fig. 2. CO₂ seasonal cycles from two years of the GMI Combo CTM. **(a)** The black line is the average of Mauna Loa and Samoa boundary conditions (derived from the NOAA Global Monitoring Division Flask Sampling Network), the dashed blue line is that cycle lagged by 2 months, and the red line is the model tropical 380 K cycle, considered to be the stratospheric boundary condition for CO₂. **(b)** The stratospheric boundary condition at 380 K (red) and the tropical cycle at 435 K (solid blue); there is a 2.5 month lag between them. The dashed blue line approximates how the model cycle would look if attenuation and lag followed the aircraft results from B96. **(c)** The stratospheric boundary condition at 380 K (red) and the CO₂ cycle in the midlatitudes, 380–400 K, having N₂O in the range of 305–310 ppb.

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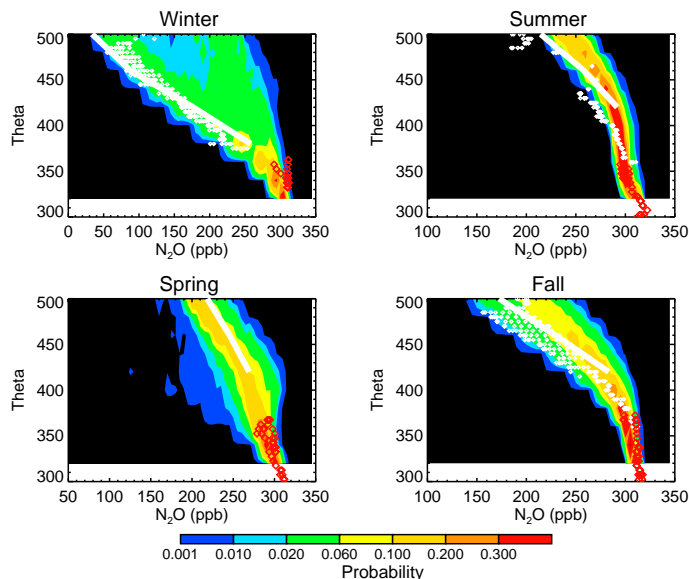


Fig. 3. Comparison of GMI N_2O profiles ($70\text{--}88^\circ\text{N}$) in four seasons with MLS, SPURT, and ER-2 data. Contoured pdfs are averages over the same seasonal sampling periods of the SPURT data (all months of the year sampled except March, June, September, and December). Yellows and reds indicate sharply peaked distributions. MLS data are shown as most probable seasonal values (solid white line). ER-2 data are seasonal means (white points) and SPURT data are seasonal averages (red points).

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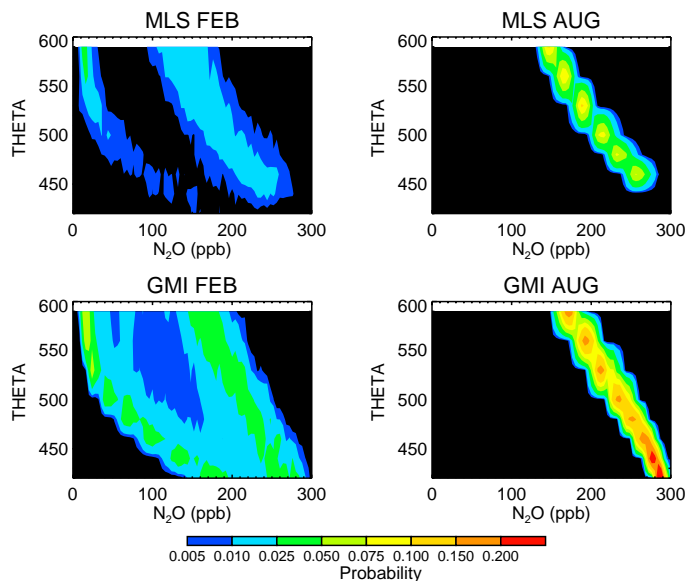


Fig. 4. Comparison of GMI and MLS N_2O profile variability (66° – 82° N). MLS data is shown for 2005 and 2006. GMI variability is averaged over 2 consecutive model years.

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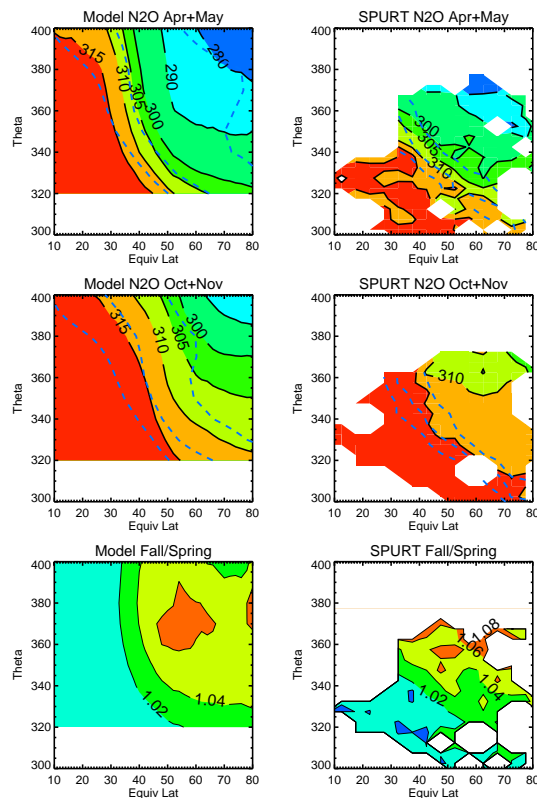


Fig. 5. N_2O from GMI and SPURT in spring and fall in the dynamical coordinate of potential temperature and equivalent latitude. Potential vorticity contours (2, 4, and 6 PVU) are overlaid with dashed blue lines. Bottom panels show the fall/spring ratio.

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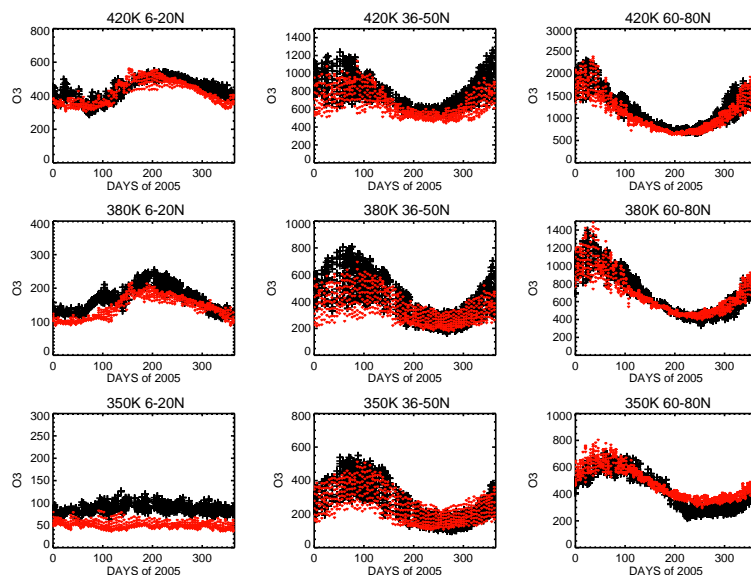


Fig. 6. Comparison of GMI (red) and MLS (black) O₃ seasonal cycles for three latitude bands and 3 levels in the UT/LS.

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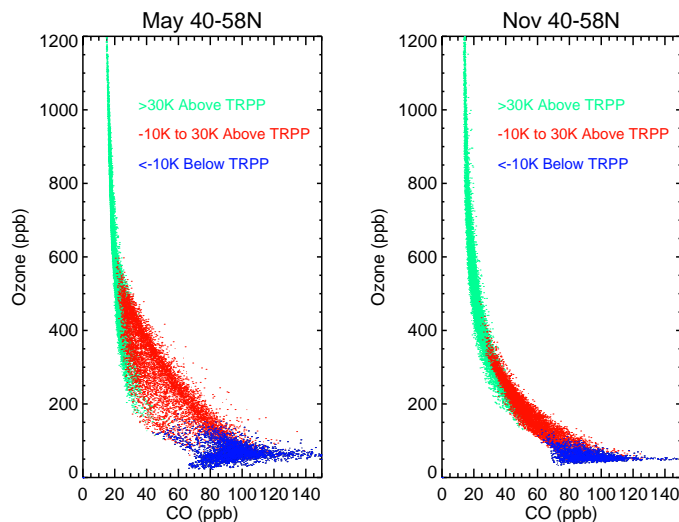


Fig. 7. Scatterplot of GMI CO and O_3 in the UT/LMS, spring and fall. Points are color-coded by their location with respect to the dynamical tropopause (2 PVU). Blue points are more than 10 K below the tropopause. Green points are more than 30 K above the tropopause. Red points, which are just below and up to 30 K above the tropopause, represent the mixed region near the tropopause.

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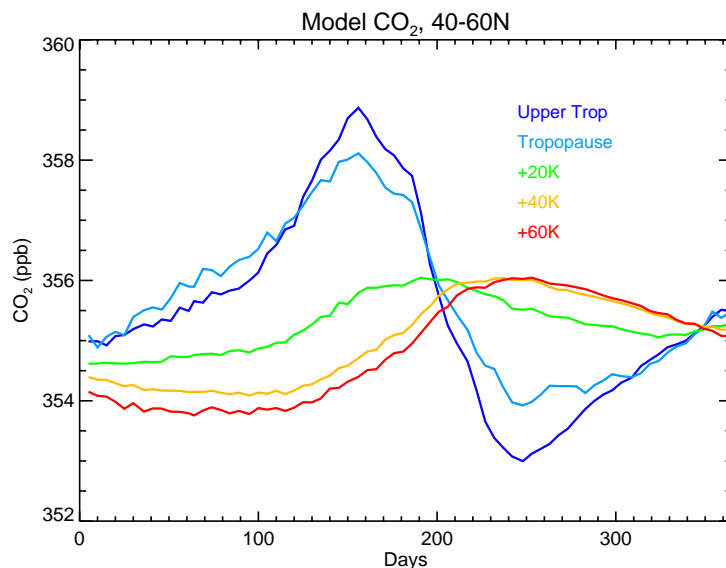


Fig. 8. GMI CO₂ cycles just below, at, and above the midlatitude tropopause. The trend in CO₂ has been removed to clarify the differences in cycle phase and amplitude across the tropopause.

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